FRUNZE, T.M.; KORSHAK, V.V.; ANDREYEV, D.N.; KUKHARSKAYA, E.V.

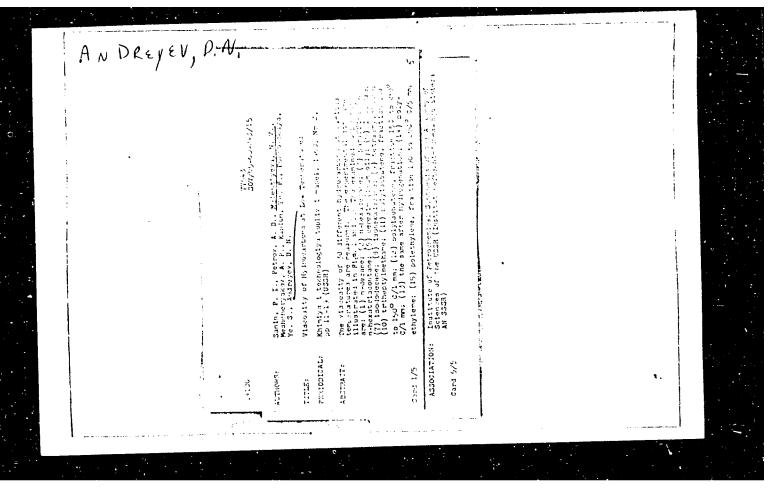
Heterochain polyamides. Part 16: Polyamides containing siloxane groups in the main chain. Vysokom.soed. 1 no.4:489-494 (MIRA 12:9)

Ap '59.

1. Institut elementoorganicheskikh soyedineniy An SSSR, 1 Institut khimii silikatov AN SSSR.

(Amides)

"APPROVED FOR RELEASE: 03/20/2001 CIA-RDP86-00513R000101510015-7



Andreyev, D.N.

820h2 \$/062/60/000/02/05/012 B003/B066

53700

AUTHOR:

Andreyev, D. N.

TITLE:

Condensation of Dimethyl-dichloro Silane in Silent

Discharges

PERIODICAL:

Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk

1960, No. 2, pp. 237 - 243

TEXT: The author condensed the dimethyl-dichloro silane in silent discharges by using a method described in previous papers (Refs. 1, 2), and investigated the composition of the resultant condensate. After and investigated the composition of the resultant condensate after fractionation of the reaction product, the individual compounds were fractionation of the reaction product, the individual compounds were identified, among other things, by converting the chlorine containing substances to silicon hydrocarbons by means of CH<sub>2</sub>MgBr, or by determining

the moral refraction. The condensate mainly consisted of polymethyl-silane chlorides, the principal chain of which possessed C- and Si-atoms in alternate succession. The yield of the condensate was 68 .. 70 per cent of the initial amount of (CH<sub>3</sub>)<sub>2</sub>SiCl<sub>2</sub>. The following compounds were

Card 1/2

DOLGOV, B.N. [deceased]; KUKHARSKAYA, E.V.; ANDREYEV, D.N.

Organosilicon esters of acrylic and methacrylic acid. Part 2:
Polymerization and properties of the polymers. Vysokom. soed. 2
no.10:1463-1465 0 '60.

1. Institut khimii silikatov AN SSSR.

(Acrylic acid) (Methacrylic acid) (Folymers)

82683

s/079/60/030/008/008/008 B004/B064

5 3700

Andreyev, D. N., Kukharskaya, E. V. AUTHORS:

Organosilicom Esters of Acrylic and Methacrylic Acid.

I. The Synthesis of Monomers TITLE:

Zhurnal obshchey khimii, 1960, Vol. 30, No. 8, PERIODICAL:

pp. 2782 - 2784

TEXT: The organosilicon esters of acrylic- and methacrylic acid were synthesized according to a procedure already described in a previous paper (Ref. 6) concerning the production of organosilicon esters of propionic and isobutyrio acid: R<sub>4-x</sub>SiCl<sub>x</sub> + xMOCOC(R')=CH<sub>2</sub>

 $\rightarrow R_{4-x} \text{Si}\left[\text{OCOC}(R^{t})=CH_{2}\right]_{x} + xMC1, (R = CH_{3} \text{ or } C_{2}H_{5}, R^{t} = H \text{ or } CH_{3},$ M = alkaline or alkaline-earth metal). The yields were 30-70%. Inhibitors (hydroquinons, picric acid, etc.) had to be added to the reaction mixture since the esters polymerize easily. The monomers with three acryl- or methacryl radicals could be isolated only by freezing at -50 to -80°C. The reaction with the free acids gave lower yields. The

Card 1/2

82683

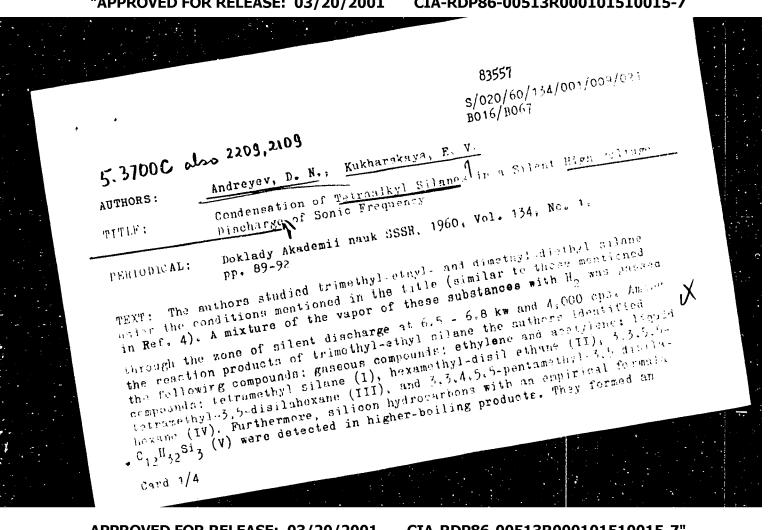
Organosilicon Esters of Acrylic and Metha- S/079/60/030/008/008/008 crylic Acid. I. The Synthesis of Monomers B004/B064

esters are colorless liquids or crystalline substances with low melting point. They hydrolyze readily. The silicon content was determined by mineralizing with a mixture of concentrated sulfuric and nitric acid, the molecular weight cryoscopically in benzene, the ester numbers by the molecular weight cryoscopically in benzene, the ester numbers by saponification with KOH in diethylene glycol. Synthesis was carried out of: trimethyl silyl acrylate, dimethyl silyl diacrylate, methyl silyl acrylate, as well as the corresponding ethyl compounds and methacrylates. A table gives the physical and analytical data. There are 1 table and 6 references: 3 Soviet, 2 US, and 1 British.

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institute of Silicate Chemistry of the Academy of Sciences USSR)

SUBMITTED: July 29, 1959

Card 2/2



CIA-RDP86-00513R000101510015-7" **APPROVED FOR RELEASE: 03/20/2001** 

### 83557

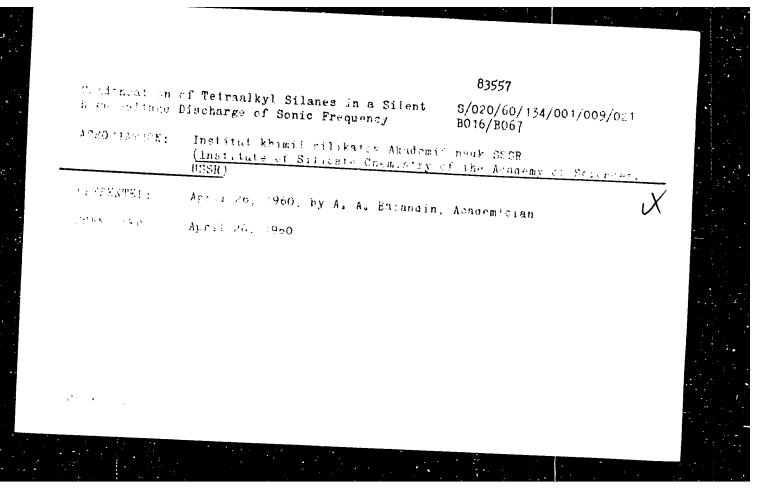
Condensation of Tetraalkyl Silanes in a Silent High-voltage Discharge of Sonic Frequency S/020/60/134/001/009/021 B016/B067

isomeric mixture difficult to separate. The high-molecular polymers with a boiling point >220°C had an average molecular weight of 1690, and corresponded to the composition (C<sub>3.8</sub>H<sub>9</sub>Si)x. The ratio C/Si in the condensation products was considerably lower than in the initial (CH<sub>3</sub>)<sub>3</sub>SiC<sub>2</sub>H<sub>5</sub>; it decreased monotonously from 5: 1 to 3.8: 1. The authors conclude theree decreased monotonously from 5: 1 to 3.8: 1. The authors conclude theree

decreased monotonously from 5: 1 to 2.5 this succeeded in bringing from that the polycondensation process which they succeeded in bringing about is no simple dehydrogenating condensation by separation of the about is no simple dehydrogenating condensation by separation of the CaH bonds but that it is much more complicated due to the runture wither of the Si-CH<sub>3</sub> or the Si-C<sub>2</sub>H<sub>5</sub> bonds. The authors proceed that the

firstently high so that they are not subject to ruptures with mentioning in the experiments described here. The authors regard the dissociation of the Si C bonds as the primary effect under the action of discharga. Further transformations of the methyle and ethyl radicals lead to the formation of C2H2, C2H4, and H. The lack of compounds with a Si Si bond in the condensate suggests that no recombination takes place among the organisation radicals. A. A. Balandin Ya. G. Eydus, and N. G. Zalegin organisation radicals. A. A. Balandin Ya. G. Eydus, and N. G. Zalegin

Card 1/4



84637

s/020/60/134/004/011/023 B016/B060

15.8114

Andreyev, D. N. and Kukharskaya, E. V.

AUTHORS:

Condensation of Hexamethyl Disiloxane in a High-voltage Dark

Discharge of Sonic Frequency

PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol. 134, No. 4,

pp. 817 - 820

TEXT: The various ways of producing organosilicon compounds with alternating silicon hydrocarbon links (-Si-(CH<sub>2</sub>)<sub>n</sub>-Si-) and siloxane links (-Si-O-Si-) are inadequate in several respects. In view of this fact, the authors thought of bringing about a condensation of hexamethyl disiloxane by means of a single passage through the zone of electric discharge. Since the compounds used in the process remain in this zone for only a short time, the compounds used in the process remain in this zone for only a short time, secondary processes were expected to be eliminated to a large extent. The secondary processes were expected to be eliminated to a large extent. The discharge tube was fed with a current of 4000 periods per sec. The procedure applied by the authors ensured good yields of low-molecular condensation products: voltage 50 kv, throughput 8 - 9 ml/h. A passage of

Card 1/3

84687

Condensation of Hexamethyl Disiloxane in a S/020/60/134/004/011/023 High-voltage Dark Discharge of Sonic Frequency E016/E060

lower-molecular polymers during interaction with H or  $\mathrm{CH}_3$ , or due to the cleavage of individual bonds in low-molecular polymers under the direct action of electric discharge. The resulting new radicals are recombined to form higher-molecular compounds. The presence of octamethyl trisiloxane in the reaction products gives evidence of a partial cleavage of the Si-O bonds. Octamethyl trisiloxane is probably the product of a recombination of two radicals (see Scheme (I)). The polycondensation carried out by the authors is accompanied by a partial but moderate demethylation (C/Si = 2.5 - 2.7, as against the hexamethyl disiloxane employed, where C/Si = 3). The method recommended here may be also used to condense other organosiloxanes. There are 1 figure and 14 references: 7 Soviet, 5 US, and

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR ( <u>Institute of Silicate Chemistry of the Academy of Sciences USSR</u>)

PRESENTED: May 4, 1960, by A. V. Topchiyev. Academician

SUBMITTED: May 4, 1960

Card 3/3

5.3700

37760 \$/661/61/000/000/021/081 D205/D302

AUTHOR:

HOR: Andreyev, D. N.

TITLE:

Condensation of methyl chlorosilanes in a silent electri-

cal discharge

SOURCE:

Khimiya i prakticheskoye primeneniye kremneorganicheskikh soyedineniy; trudy konferentsii, no. 6, Doklady, diskussii resheniye. 11 Vses. Konfer. po khimii i prakt. prim. kremneorg. soyed., Len. 1958. Leningrad. Izd-vo AN SSSR.

1961, 109

TEXT: The behavior of SiCl<sub>4</sub> (in H<sub>2</sub> medium) under the influence of a silent discharge at 24 - 25 kV and 25 c/s was investigated earlier, showing that the SiCl<sub>4</sub> does not suffer changes in these conditions. Recently, in discharges of a 4000 c/s frequency, it was found that SiCl<sub>4</sub> vapors are condensed with the formation of a mixture of roughly equal amounts of Cl<sub>3</sub>SiSiCl<sub>3</sub> and Cl<sub>3</sub>SiSiCl<sub>2</sub>.SiCl<sub>3</sub>.

Card 1/2

X

Condensation of methyl ... S/661/61/300/006/021/051
The yield of the octachlorotrisilane is 10% calculated on the initial SiCl<sub>4</sub> and 40 - 45% on the reacted SiCl<sub>4</sub>.

ASSOCIATION: Institut khimii silikatov, AN SSSR, Leningrad (Institute of Silicate Chemistry, AS USSR, Leningrad)

25369 -S/079/61/031/008/007/009 5 3700 D215/D304 AUTHORS: Andrewev, D.N., and Lyutyy, V.P. TITLE: Synthesis and properties of silicon organic complex PERIODICAL: Zhurnal obshchey khimii, v. 31, no. 8, 1961, 2726-2729 TEXT: The authors undertook this experimental work on the supposition that these complex silicon compounds could be as useful as similar normal organic ones. They obtained nine new esters from alkali salts of higher aliphatic and aromatic acids and halogen derivatives of alkyl--silanes, and siloxanes by heating the components in dimethylformamide, according to the following schemes  $2 \mbox{${\rm KCOOK}$} + \mbox{${\rm CICH_2Si(CH_3)_2O(CH_3)_2SiCH_2Cl} \longrightarrow$ The constants of obtained diesters are given in the Table.  $\rightarrow RCOOCH_2Si(CH_3)_2O(CH_3)_2SiCH_2OCOR + 2KCl$ 2(CH<sub>3</sub>)<sub>3</sub>SiCH<sub>2</sub>Cl + KOCOR'COOK → (see next card) → (CH<sub>3</sub>)<sub>3</sub>SiCH<sub>2</sub>OCOR'COOCH<sub>2</sub>Si(CH<sub>3</sub>)<sub>3</sub> + 2KCI Card 1/4

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Syn	Synthesis and properties				25369 .S/079/61/031/008/007/009 D215/D304							
(1)K	оистанты	полученных диа	фироп пред	(стаплон	i o ti	пблице	9.				`	
M n.n.	(3)	Формула ,	Температура кипения (дан ление в мм)	- CTSIBBURG	. 20	d.**	ияя- дено	пычн- слено		•		
(I) (III) .	(CII3CII2C (CII3)2CII (C <sub>0</sub> II <sub>5</sub> CO	OCH*2!(CH*)*]*0 COOCH*2!(CH*)*]*0 OCH*2!(CH*)*]*0	143° (4) 142 (2) 231 (2)	1-29° 111080-71 -45		0.9850 0.9632 1.0905	79.63 88.80 110.67	80.24 89.50 110.20		•		
(IV) (VI) (VII)	(CII <sub>3</sub> ) <sub>3</sub> [(CII <sub>3</sub> ) <sub>3</sub> SiC (CII <sub>3</sub> ) <sub>3</sub> SiC	SICH <sub>2</sub> OOCCH <sub>2</sub> ] <sub>2</sub> CH <sub>2</sub> OOC(CH <sub>2</sub> ) <sub>3</sub> ] <sub>2</sub> CH <sub>2</sub>		(отекло) 23 26 70 (стекло)	1.4382 1.4419 1.4446	0.9548 0.9497 0.9325	79.90 88.74 102.85	80.49 89.75 103.64		<b>!</b> :	•	
(VIII) (IX)	[(CII <sub>3</sub> ) <sub>3</sub> S [(CII <sub>3</sub> ) <sub>3</sub> Si	SICH <sub>2</sub> OOC(CH <sub>2</sub> ) <sub>4</sub> ] <sub>2</sub> ICH <sub>2</sub> OOC  <sub>2</sub> C <sub>6</sub> H <sub>4</sub> -0 ICH <sub>2</sub> OOC  <sub>2</sub> C <sub>6</sub> H <sub>4</sub> -0	201—202 (6) 185 (7)	-28  115-116	1.4443 1.4973 —	0.9261 1.0277 —	107.53 96.44	108.27 95.94			•	
Card	2/4	(For L	egend of	the Ta	h lo	500	4		1.		,	ħ

	-25369	
Synthesis and properties  Legend of the Table:	S/079/61/031/008/007/009 D215/D304	
(1) Constants of the obtained diest (2) N N (3) Formula (4) b.p. (pressure in mm)  (8) MRD found/calculated.	(5) temp as (c) as	
They observed that Al	"" in the series:	

25369 8/079/61/031/008/007/039 D215/D304

Synthesis and properties...

became overcooled, crystallizing afterwards spontaneously. For these esters (N I, IV, V and VII) approximate m.p. were evaluated from cooling curves. Diesters with even carbon numbers crystallize in a narrow temperature range: from -23 to -28° C. The azelaic acid ester sets at -70° C to a glassy substance; among diesters with soloxanic bond onyl NI crystallizes by cooling, N III solidifying to a glassy substance, and the di-isobutyrate (N II) stays liquid even at -70 C. There are 1 table and 10 references: 3 Soviet-bloc and 7 non-Soviet-bloc. The references to the English-language publications read as follows: -T.L. Speier, B.F. Daubert, R.R. Gregor, Y.Am.Chem. Soc. 71, 1474, (1949); R.L. Merker, T.E. Noll, Y. Org. Ch. 21, 1537, (1956); L.H. Sommer, W.D. English, G.R. Ansul, D.N. Vivona, S.Am. Chem. Soc. 77, 2485 (1955); L.H. Sommer, N.S. Marans, G.M. Godberg and others, S. Am. Chem. Soc. 73, 882, (1951).

ASSOCIATION: Institut khimii silikatov akademii nauk SSSR (Institute of

Silicates, Academy of  $S_{C}$ iences USSR)

SUBMITTED:

August 5, 1960

Card 4/4

15 7140

27067 S/080/61/034/003/007/017 A057/A129

AUTHORS:

Andreyev, D. N., Okhrimenko, I. S., Pinchuk, A. Ye., Lyutyy, V. P.

TITLE:

Unsaturated organisilicon polyesters and the properties of lacquers

PERIODICAL: Zhurnal prikladnov khizii, v. 34. no. 3, 1961, 584 - 588

TEXT: Syntheses of two new polynumarates, modified with siloxane links, viz., the diester bis(trimethylsilylmethyl)fumarate and the polyester poly(dimethylenetetramechyldisiloxane)fumarate are described and preliminary results in investigations of their properties are given. Polymaleates and polyfumarates are widely used in the manufacture of lacquers, plastics etc. M. A. Bulatov and S. S. Spasskiy [Ref. 1: Vysokomol. soyed., 2, 5, 658 (1960)] demonstrated already that these esters, when modified with organosiloxanes, as for instance with dimethyldiethoxysilane, obtain high technical properties. Organosiloxane polymaleates and polyfumarates, derivatives of organosiloxane and glycols containing a siloxane link in the molecule, hav. not been synthesized. Thus the present authors started investigations in this field. To develop the synthesis of the polyester, the synthesis of the diester was carried out first according to the reaction 2(CH3)3SiCH2Cl +

Card 1/5

Unsaturated organosilicon polyesters and the...

27067 \$/080/61/034/c03/007/017 A057/A129

+ KOOCCH=CHCOOK → (CH3)3S1CH2OOCCH=CHCOCCH2S1(CH3)3. In the procedure 0.16 mole of potassium fumarate was mixed with 0.08 mole of fumaric acid in 150 ml of dimethylformamide as solvent. The mixture was boiled, agitated during 30 minutes, dropwise 0.32 mole chloromethyltrimethylsilane was added, heated, agitated for 20 hrs more, while the boiling temperature rose from 124°C to 149°C. After cooling the precipitated potassium chloride was filtered off, the solvent was removed by vacuum distillation and the residue was treated with a sods solution and water-benzens mixture. Then the benzene was dried, evaporated and the fumarate was vacuum-distilled (1510-153°C, 8 torr). The yield was 54.8% of a product with np 1.4548, df 0.985. In an analogous way the polyester was prepared. Potassium furarate of C15 mole was mixed with 0.125 mole of fumaric acid in 300 ml of dimethylformamide and then bis(cnloromethyl)tetramethyldisilexane was added. Instead of benzene ether was used as solvent and after removal of the latter a highly viscous reddish-brown sutstance insoluble in water but soluble in most organic solvents, except petroleum ether and gasoline, was obtained. The average molecular weight was found to be 2,400 corresponding to a condensation degree of 9. The re-precipitated polyester was investigated by spectrophotometry on an NKC-12 (IKS-12) apparatus. The obtained infrared absorption spectrum proved the presence of double bonds in the trans-

Card 2/5

Unsaturated organosilicon polyesters and the...

27067 8/080/61/034/003/007/017 A057/A129

position (900 - 990 cm<sup>-1</sup>, 1,320 cm<sup>-1</sup>), siloxane bonds (1,020 - 1,091 cm<sup>-1</sup>), (CH<sub>3</sub>)<sub>2</sub>Si groups (800 - 814 cm<sup>-1</sup>, and 1,259 cm<sup>-1</sup>), ester groups characteristic for fumarates (1,140 - 1,180 cm<sup>-1</sup>) and end-carboxylic groups (900 - 950 cm<sup>-1</sup>). The obtained polyester is miscible with styrene within a range from 3.5: 1 to 0.3: 1, and transparent homogeneous solutions are obtained. Properties of four lacquers (see table) hased on this polyester were investigated and it was observed that in comparison with maleic resins the double bonds in siloxane-modified fumaric resin show lower activity. Thus a lacquer based on this resin required heating to 200°C to "dry" within 30 minutes, i.e., to produce a three-dimensional structuration to 70% (Fig. 3). At 120°C the same degree of structuration requires 7 hrs (Fig. 2). The drying is the result of two independent processes: 1) a catalytic polymerization with an initiator (1% of cyclohexanone peroxide), and 2) an oxidative structuration produced by heating over 100°C. No initiator seems to be necessary for the latter process. Elasticity tests carried out by the NIILK method and hardness tests on a CMNM-4 (GIPI-4) machine showed that films obtained from these lacquers have high elasticity, but rather low hardness. Thus lacquer no. III showed after 70 minutes of drying at 200°C a 1 mm flexibility on a rod and a 0.195 hardness. There are 3 figures, 1 table, and 2 Soviet-bloc references. SUPMITTED: July 9, 1960

Card 3/5

360:4 \$/079/62/032/004/005/010 D204/D301

AUTHORS:

Andreyev, D.N., Dolgov, B.N. (Deceased) and Butts, S.V.

TITLE:

Stability of the Si-R bonds in \u03c4-silico-organic acids

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 4, 1962, 1275-1277

TEXT: The action of conc.  $H_2SO_4$  on methyl-di(n-amyl)-silyl-propionic acid showed that the Si-2n-4 Am bond is slightly more stable then the Si-180-Am bonds studied in an earlier work. Complete fission of this bond occurred on heating with  $H_2SO_4$  to 700C, over 1 hr. and keeping it for a further hour at that temperature, with stirring, to give  $[HOOC.CH_2CH_2Si(n-C_5H_1)CH_3]_2O$ , (A). Including earlier results, the authors therefore concluded that stability of Si-C towards conc.  $H_2SO_4$  in acids of the general formula  $R_2(CH_3)SiCH_2CH_2$  COOH decreases in the order Et, n-Pr > Me > n-Am > n-Bu, 180-Am. Synthesis of  $CH_3(PhCH_2)_2SiCH_2CH_2COOH$  was attempted to find whether the  $Si-CH_2X$  bond would be as stabilized by substituting Ph into the

Card 1/2

Stability of the Si-R bonds in ...

S/079/62/032/004/005/010 D204/D301

CH<sub>3</sub> group as it is by substitution with halogens, but both benzyl groups were found to break off the Si during alkaline hydrolysis of the methyl-dibenzyl-silyl-methyl malonate. Preparation of new compounds CH<sub>3</sub>(n-C<sub>5</sub>H<sub>11</sub>)<sub>2</sub>SiCH<sub>2</sub>Cl, CH<sub>3</sub>(PhCH<sub>2</sub>)<sub>2</sub>SiCH<sub>2</sub>Cl, CH<sub>3</sub>(n-C<sub>5</sub>H<sub>11</sub>)<sub>2</sub>SiCH<sub>2</sub>CH(COOEt)<sub>2</sub>, CH<sub>3</sub>(n-C<sub>5</sub>H<sub>11</sub>)<sub>2</sub>SiCH<sub>2</sub>CH<sub>2</sub>COOH and A is described and their physical properties are tabulated. There are 1table and 4 references: 3 Soviet-bloc and 1 non-Soviet-follows: L.H. Sommer, W.P. Barie, and J. Gould, J. Am. Chem. Soc., 75, 3765, 1953.

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institute of Silicate Chemistry of the Academy of Sciences, USSR)

SUBMITTED: April 19, 1960

Card 2/2

16.6173

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5/079/62/032/004/009/010 D287/D301

AUTHORS:

Andreyev. D.N., and Kukharskaya, E.V.

TITLE:

Organosilicon diethers and diesters of organosilicon

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 4, 1962, 1353-1354

TEXT: The authors found that the title compounds can be readily prepared by condensing the K or Na salts of γ-organosilicon acids prepared by condensing the k or Ha Balto of policial street with R SiCl, mono- and dibasic γ-organosilicon acids enter into this reaction. The condensation reaction was carried out by heating the components in hydrocarbon solvents. [(CH<sub>3</sub>)<sub>3</sub>SiCH<sub>2</sub>CH<sub>2</sub>COO]<sub>2</sub>Si(CH<sub>3</sub>)<sub>2</sub> (yiela 59 %) and [(CH3)3SiOCOCH2CH2Si(CH3)2]20 (Yiela: 36 %) were prepared by this method. Physical constants of the 2 compounds are given. There are 2 references: 1 Soviet-bloc and 1 non-Soviet-bloc. The reference to the English-language publication reads as follows: A.G. Brook, J. Am. Chem. Soc., v. 77, 1955, 4827.

Card 1/2

Organosilicon diethers and ...

S/079/62/032/004/009/010 D287/D301

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institute for Silicate Chemistry of the AS USSR)

SUBMITTED:

December 22, 1961

Card 2/2

S/079/62/032/004/008/010 D287/D301

AUTHORS:

Andreyev, D.M., and Kukharskaya, E.V.

TITLE:

Interaction of organomagnesium complexes with

organosilicon esters

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 4, 1962, 1352-1353

TEXT: The present work was carried out because of the absence of information on the interaction between organosilicon esters and Grignard reagents, especially between compounds of the formula RMgX with alkyl acyloxysilanes (R'Si(OCOR'')<sub>4-n</sub> where n = 1-3. The above compounds were found to form tertiary alcohols on reaction with C<sub>2</sub>H<sub>5</sub>MgBr as proved for trimethylsilyl propionate (I) and diethylsilyl dipropionate (II). Both compounds were synthesized in the usual way. In the first case 68 g of compound I was added to the Grignard reagent; the yield of 3-ethyl-3-pentanol was 32.9 %. This compound reagent; the yield of 3-ethyl-2-pentene by heating with anhydrous could be dehydrated to 3-ethyl-2-pentene by heating with anhydrous CuSo<sub>4</sub>. In the second experiment 84 g of compound II was added to

Card 1/2

Interaction of or (anomagnesium ...

\$/079/62/032/004/008/010 D287/D301

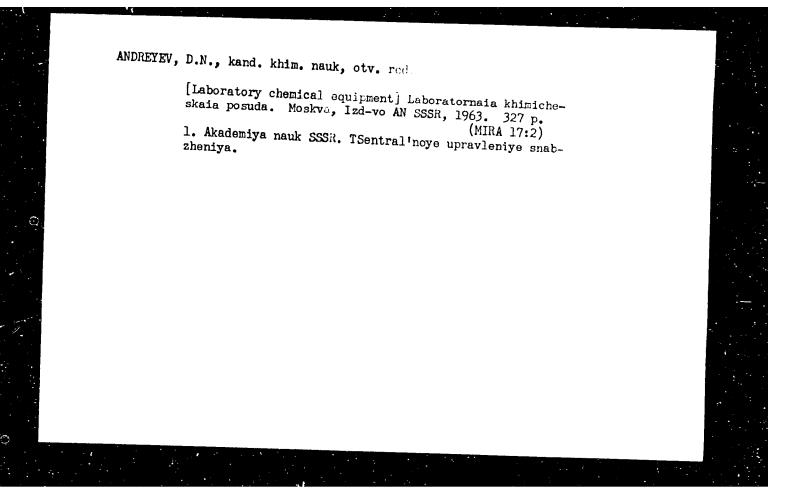
the Grignard reagent; the alcohol was obtained with 42.5 % yield. Physical constants of the synthesized compounds are given and compared with literature data. There are 3 non-Soviet-bloc references: The references to the English-longuage publications read as follows: A.A. Morton and J.R. Stevens, J. Am. Chem. Soc., v. 53, 1951, 2244; G. Egloff, Physical Constants of Hydrocarbons, v. 1, 1939, 207; M. Mtienne, C.T., 235, 967, 1952.

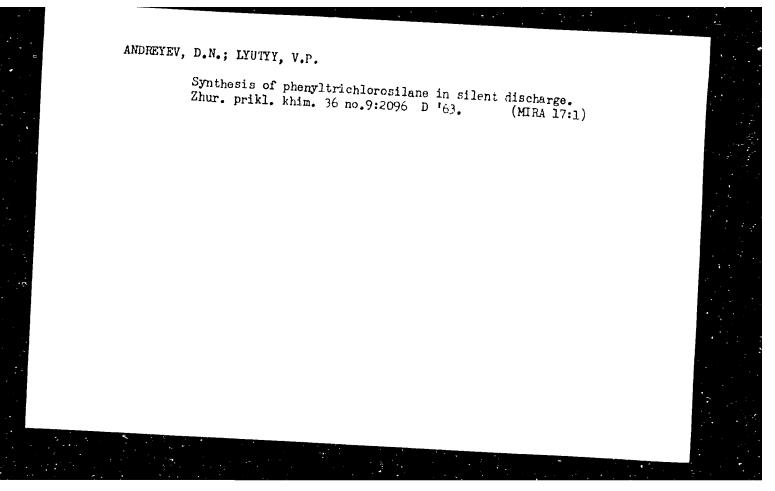
ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institu-

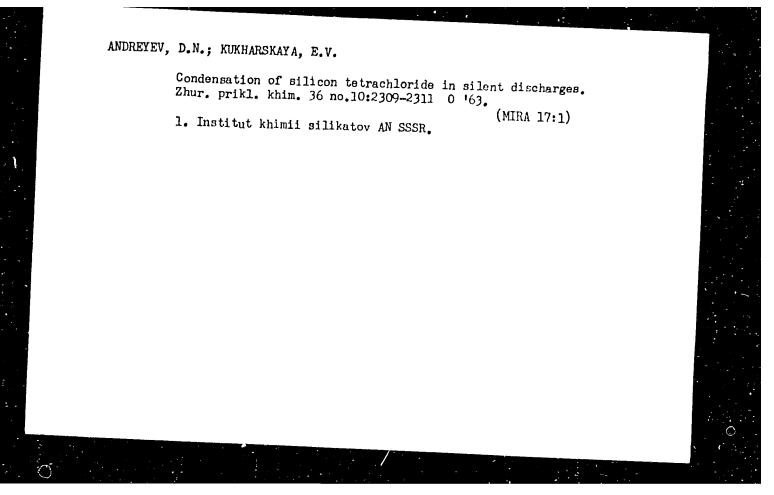
te for Silicate Chemistry of the Academy of Sciences of the USDR)

SUBMITTED: December 22, 1961

Card 2/2

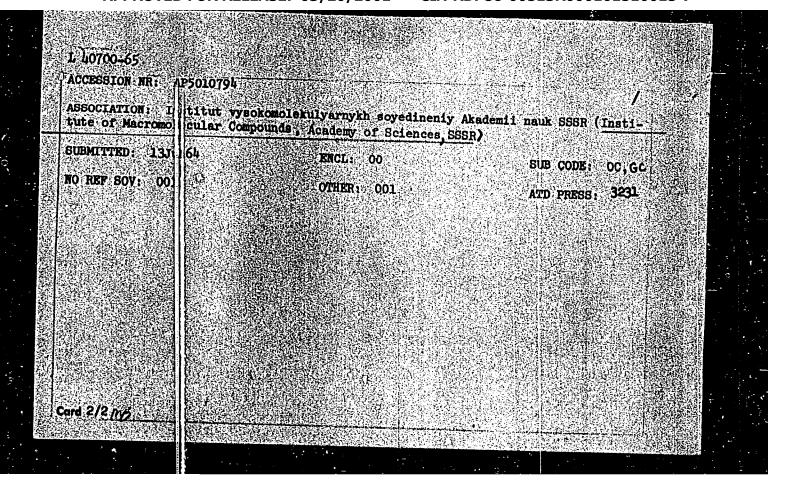






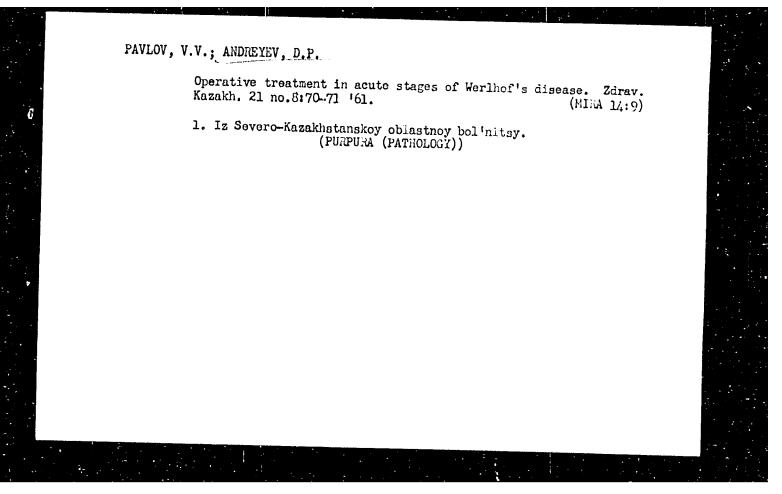
ACCESSIO	ON MR: "AP5010794	( <b>a</b> )/1 Po-1/Pr-1	UR/0079/65/035/0	04/0756/0756	<i>V.</i> "
AUTHOR:	Andreyev, D. N.; Usa	cheva, N. T.		23	
7.5		ethacryloxypolydimethyle	是不是是一个人的。 第二章	7	
SOURCE:	Zhurnal obshchey khi	mii, v. 35, no. 4, 1965	756		
TOPIC TA	AGS: polymer, polydim	ethylsiloxane, siloxane,	methacrylate.copo	lymer	
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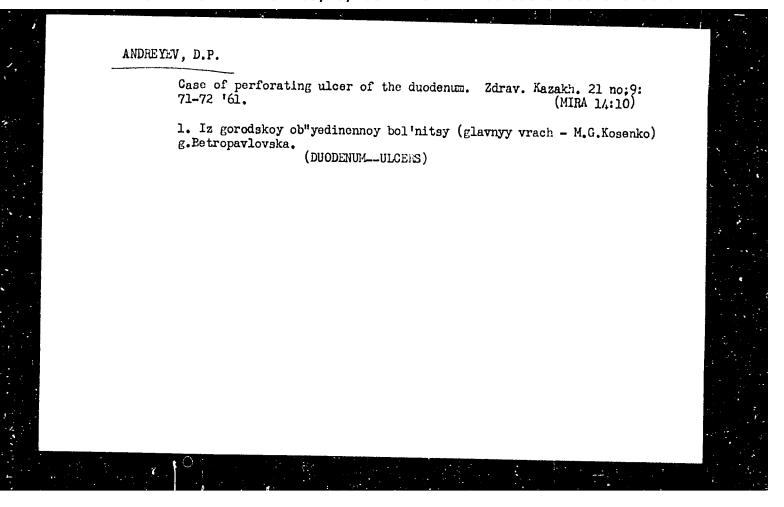
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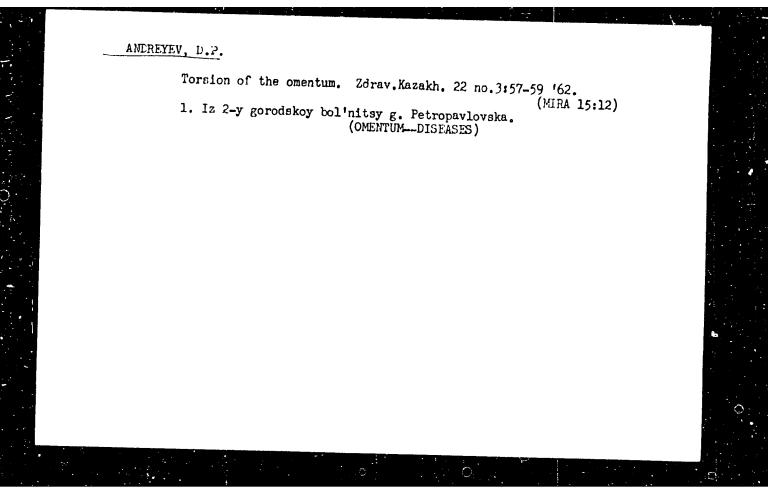


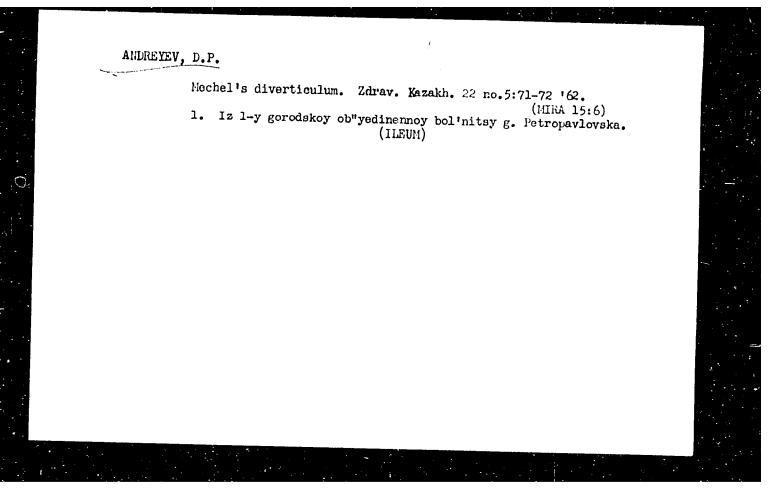
L 44564-66 EWT(m)/EWP(j) ACC NRI AP6031381 SOURCE CODE: UR/0079/66/036/009/1628/1633 AUTHOR: Amdreyev, D. N.; Afanas'yeva, G. S. 24 ORG: Institute of Macromolecular Compounds, Academy of Sciences, SSSR (Institut vysokomolekulyarnykh soyedineniy Akademii nauk SSSR) Organosilicon polyamides. I. Synthesis of dicarboxylic aliphatic organosilicon TITLE: acids SOURCE: Zhurnal obshchey khimii, v. 36, no. 9, 1966, 1628-1633 TOPIC TAGS: organosilicon compound, organosilicon polyamide, organosilicon ABSTRACT: This article is the first in a series on the study of organosilicon polyamides. Some polyamides from dicarboxylic acids which contain siloxane links in their chain are known. Polyamides from a dicarboxylic acid with one silicon atom in its chain are mentioned in the literature, but not described. Synthesis of two siliconcontaining dicarboxylic acids was prompted by the desire to study the effect of the length of the acid chain on the properties of the polyamides obtained from such acids. The general formula of these acids was  $(CH_3)_2Si/(CH_2)_1COOH/_2$ . The first of the synthesized acids, with n=2, namely, 4,4-dimethyl-4-silaheptane-1,7-dicarboxylic Card 1/2 UDC: 547.245+547.461

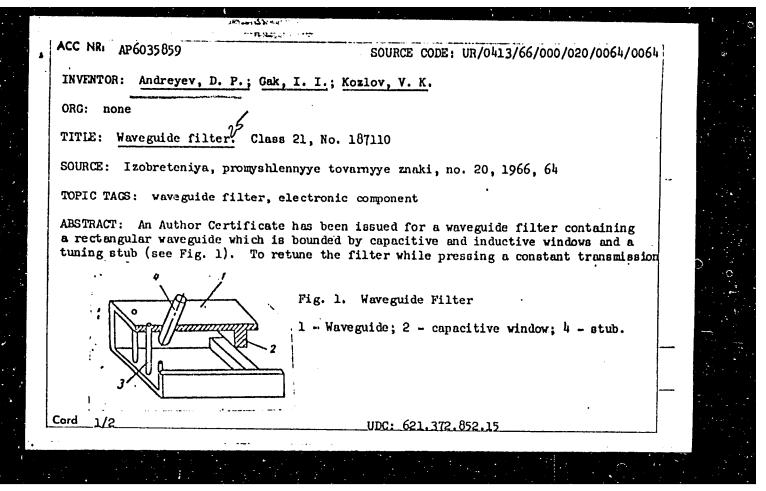
ACC NR: AP6031381		
	0	(
acid was prepared according to the available literature data. Another acid	i	
: ), ) = dimetry i = ) = Blianonane = 1,9 = dicarboxylic acid was obtained by a new method		
(1) $(CH_3)_2SiCl_2 + 2HO(CH_2)_4Cl \rightarrow (CH_3)_2Si[O(CH_2)_4Cl]_2$ .	į	
177	1	
$\frac{2(CH_{5})_{5}SiO(CH_{2})_{4}Cl}{(VIII)} = \frac{2Mg}{2(CH_{5})_{5}SiO(CH_{2})_{4}MgCl} = \frac{(CH_{5})_{5}SiO_{4}}{(CH_{5})_{5}SiO_{4}}$		
(2) $(CH_3)_2Si[(CH_2)_4OSi(CH_3)_3]_2 \xrightarrow{H_1O} (CH_3)_2Si[(CH_2)_4OH]_2$	į	
(3) (VI) $\frac{\text{KMnO}_4}{\text{NaOII}}$ (CII <sub>3</sub> ) <sub>2</sub> Si[CII <sub>2</sub> CII <sub>2</sub> CII <sub>2</sub> COOII] <sub>2</sub>	ļ	
(IX)		
The yield of the final stage was 38%. The structure of the final product and or intermediates was confirmed by ID.		
intermediates was confirmed by IR spectra. Orig. art. has: 6 formulas, 1 table	f the	
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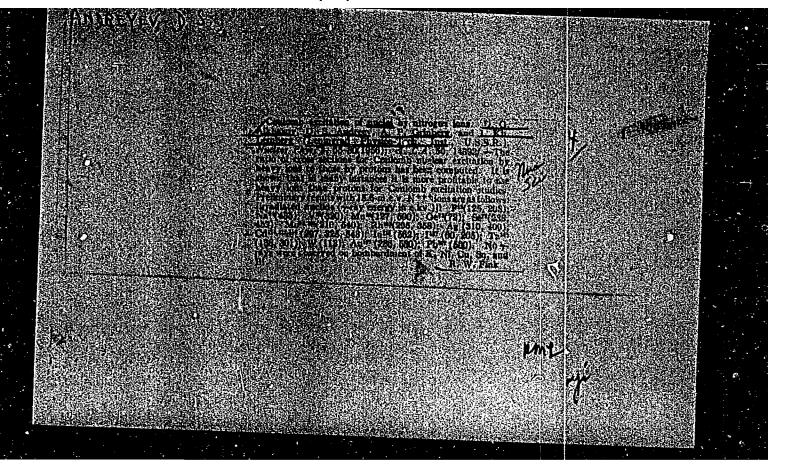




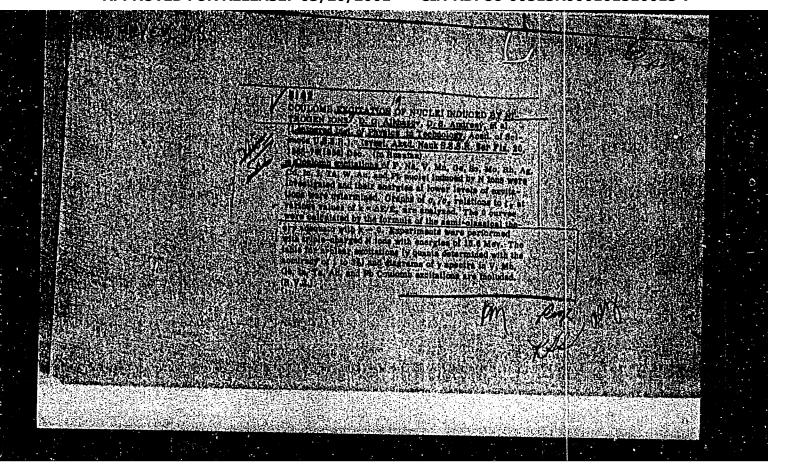


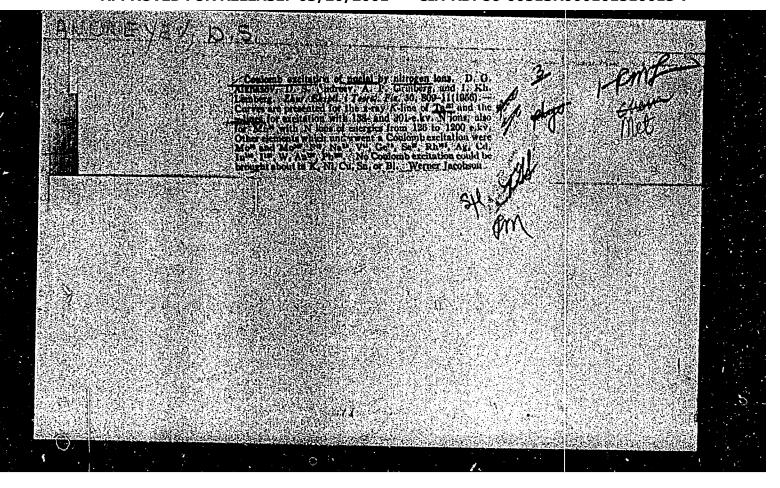
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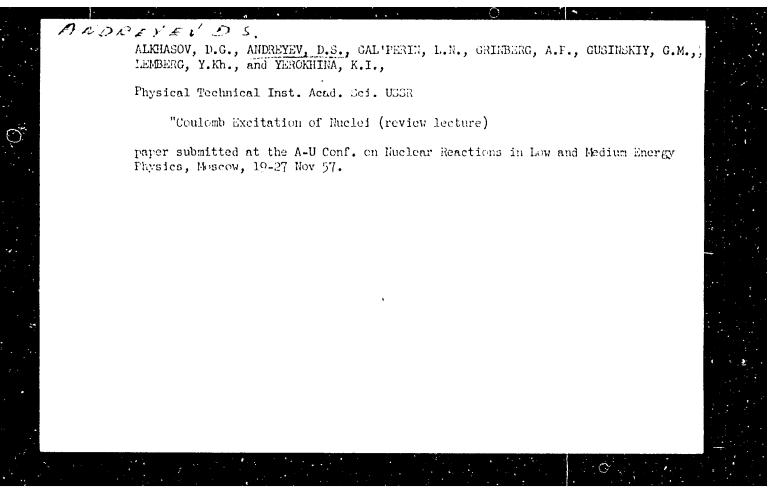
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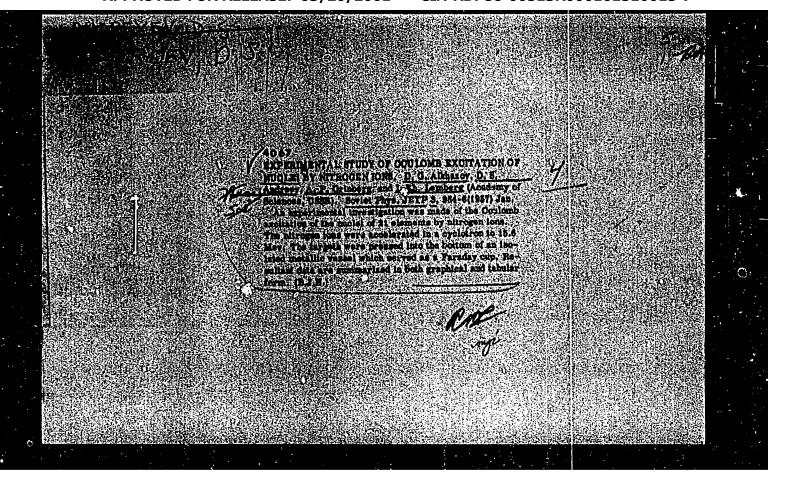


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ANDREYEV, D.S.

AU THORS:

Alkhazov, D. G., Andreyev, D. S., Yerokhina, K. I., Lemberg, I. Kh.

56-6-6/47

TITLE:

The Coulomb Excitation of Separated Tin Isotopes (Kulonovskoye vozbuzhdeniye razdelennykh izotopov

olova).

PERIODICAL:

Zhurnal Eksperimental noy i Teoreticheskoy Fiziki, 1957,

Vol. 33, Nr 6, pp. 1347-1358 (USSR)

ABSTRACT:

A 14,5 MeV  $\alpha$ -beam coming from a cyclotron is focused by a system of quadrupol lenses in an aluminum tube and thus impinges upon a target, where Coulomb excitation takes

place. The y-quanta liberated on this occasion are measured in a well screened n-scintillation spectrometer.
Between the crystal and the target the following n-absorbers

are connected:

400 μCu; 1.3 mm Al; 100 μ mica; 1 mm MgO; 50 μ Pb

The following measuring and computation results were and 1.5 mm air.

obtained:

Card 1/3

"APPROVED FOR RELEASE: 03/20/2001 CIA-RDP86-00513R000101510015-7

he Coulomb E	xcitation	of Separated Ti	n Isotopes	56-6-6/47	
	isotope	ΔE in MeV <sup>x</sup> )	$\frac{B(E2)}{e^2}.10^{48} \text{ in cm}^4$	τ.10 <sup>13</sup> in sec.	
	Sn <sup>112</sup>	1,26	0,18	7,2	
	Sn 114	1,30	0,20	5,5	
	Sn 116	1,29	0,19	6,0	
	5n 5n 118	1,22	0,19	8,0	
	Sn <sup>120</sup>	1,18	0,17	10,5	
	Sn <sup>122</sup>	1,15	0,15	13,5	
	Sn 124	1,13	0,14	15,9	
	Sn 115	_xx)	-		
	Sn 117	( 0,865 ( 1,03	(0,025 (0,09	<del>-</del>	
	Sn <sup>119</sup>	0,907	0,11	-	١.
ard 2/3	Sn <sup>117</sup>			- -	

The Coulomb Excitation of Separated Tin Isotopes

56-6-6/47

x) Corresponds to the energy of the \*-quantum which goes directly to the basic state.

xx)Between 0,75 up to 1,75 MeV no requanta were found.

There are 7 figures, 1 table, and 19 references, 4 of which are Slavic.

ASSOCIATION: Leningrad Physico-Technical Institute AN USSR

(Leningradskiy fiziko-tekhnicheskiy institut Akademii

nauk SSSR).

SUBMITTED:

June 3, 1957 (initially) and October 5, 1957 (after

revision)

AVAILABLE:

Library of Congress

Card 3/3

ANDREYEV, D. S.: Master Phys-Math Sci (diss) -- "Investigation of the coulomb excitation of dilute lead isotopes". Leningrad, 1958. 1h pm (Phys-Tech Inst Acad Sci USSR), 150 copies (KL, No 5, 1959, 183)

S/048/60/024/012/005/011 B019/B056

AUTHORS: Andreyev, D. S., Yerokhina, K. I., and Lemberg, I. Kh.

TITLE: Cascade Excitation of the Second Rotational Levels in

Separated Tungsten Isotopes 19

PERIODICAL: Izvostiya Akademii nauk SSSR. Seriya fizichenkaya, 1960, Vol. 24, No. 12, pp. 1470-1475

TEXT: The present paper was read at the 10th All-Union Conference on Nuclear Spectroscopy, which was held in Moscow from January 19 to January 27, 1960. By means of Coulomb excitation it was hitherto possible to excite even-even nuclei to the first rotational level. The excitation to the secon rotational level was not possible. Protons and &-particles were used for excitation. In the introduction, the authors discuss several results obtained by earlier papers. The experiments described here were carried out by means of quadruply-charged Nevi ions having an energy of 27.8 Mev, and by means of quadruply-charged N20 from with an energy of 36 Mev. Four metallic tungsten targets were investigated, of which the

Card 1/2

Cascade Excitation of the Second Rotational Levels in Separated Tungsten Isotopes

S/048/60/024/012/005/011 B019/B056

first consisted of 87.6% W<sup>182</sup>, the second of 90.1% W<sup>183</sup>, the third of 91.3% W<sup>184</sup>, and the fourth of 96% W<sup>185</sup>. The Maspectrum, which was emitted during the bombardment of the targets with heavy ions, was investigated by the author by means of a scintillation Maspectrometer. The photomultiplier pulses were amplified and recorded with a 63-channel amplitude analyzer, which had been developed by LETI. From the experimental results represented in diagrams and a detailed discussion, the authors became convinced that the Malines with the energies of 250 keV. 250 keV and 275 keV, which they had observed, represent the result of a double Coulomb excitation of states with the energies of 550. 360 and 400 keV in W<sup>182</sup>, W<sup>184</sup> and W<sup>186</sup> Similar results obtained by Newton et al. (Ref. 6) are mentioned. There are 3 figures 1 table, and 6 references: 2 Soviet. 2 French, and 2 US.

ASSOCIATION: Fiziko-tekhnicheskiy institut Akademii nauk SSSR (Institute of Physics and Technology of the Academy of Sciences USSR)

Card 2/2

S/046/60/024/012/006/011 BO19/BO56

Andreyev, D. S., Grinberg, A. P., Gusinskiy, G. Yerokhina, K.I., and Lemberg, I. Kh. AUTHORS:

Coulomb Excitation of the First Nuclear Levels of Even

Chromium-Selenium and Neodymium Isotopes

Izvestiya Akademii nauk SSSR. Seriya fizicneskaya, 1960, PERIODICAL:

Vol. 24, No. 12, pp. 1474-1477

TEXT: The present paper was read at the 10th All-Union Conference on Nuclear Spectroscopy, which was held in Moscow from January 19 to January 27, 1960. The experiments described in the present paper were carried out with 16.3-Mev and 36.0-Mev nitrogen ions and 23.2-Mev neon ions. Results are given in Table 1. Chromic oxide targets were used, which contain the isotopes  $\operatorname{Cr}^{52}$  and  $\operatorname{Cr}^{54}$ , as well as natural, vaporized chromium. Further, natural metallic neodymium (23.87% Nd  $^{144}$ ) was used. The results

are discussed in great detail and compared with earlier results. There are 3 figures, 1 table, and 17 references: 3 Soviet and 14 US.

Card 1/4

TITLE:

	• • •				S./04 B019	8/60/024/ /B056	/012/006/011	
1	Ž	3	4	Репериь	ц уровень		}	
 Идро	Δ E,Mev	Бомбардирующие частицы и из впергия. МеV	<b>Q</b> ядро	b <sub>ΔE, MeV</sub>	EB(E2) 1044, e2	с/ литература		
Cr <sup>50</sup> Cr <sup>52</sup> Cr <sup>51</sup> Se <sup>76</sup>	0,78 1,45 0,84 9,560	No <sup>20</sup> (23,2) N <sup>14</sup> (36,0) N <sup>14</sup> (16,3), No <sup>20</sup> (23,2) N <sup>14</sup> (36,0), No <sup>20</sup> (23,2)	Mo <sup>100</sup> . Ni <sup>58</sup> . Ti <sup>48</sup>	0,53 1,45 0,99	0,614 0,080 0,070	[3] · [1] [4]		
Se <sup>78</sup> Se <sup>80</sup>	0,615 0,650	N <sup>14</sup> (36,0), Ne <sup>20</sup> (23,2) N <sup>14</sup> (36,0)	} { Mo100	0,53 0,99	0,614 0,070	[3] [4]	<u>V</u>	
Se <sup>52</sup> Nd <sup>144</sup> Nd <sup>146</sup> Nd <sup>148</sup> Nd <sup>150</sup>	0,660 0,695 0,455 0,300 0,130	N14(36,0) N14(36,0) N14(36,0) N14(36,0) N14(36,0)	Zr°2,04 Zr°2,04 Rh103 Tn161	0,92 0,92 0,30 0,136	0,079 0,079 0,21 2,04	[3] [3] [5] [6]		
Car	rd 3/4					•		
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		5 B(E2) † .10".   C B(E2) † .10".   C  0,15 0,062 0,057 0,42   0,43 [7] 0,23   0,36 [7] 0,23   0,23 [7] 0,23 0,19 0,23 0,25   0,25 [9] 0,57   0,69 [9] 1,92   2,3 [9]	0,9 1,2 1,7 1,8 1,3 1,5 1,1	\$\frac{\fint}{\frac{\fir\fir\fir\fir\fir\fir\f{\frac{\frac{\frac{\frac{\frac{\frac{\fir}}}{\firac{\fir}{\firin}}}}}{\firac{\frac{\fir}{\firin}}}}{\fin}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}	`		
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5/048/60/024/012/007/011 BO19/BO56

Andreyev, D. S., Yerokhina, K. I., and Lemberg, I. Kh. AUTHORS:

The Coulomb Excitation of the Ne 21 Nucleus TITLE:

Izvestiya Akademii nauk SSSR Seriya fizicheskaya, 1960, PERIODICAL: Vol. 24. No. 12, pp. 1478-1479

TEXT: The present paper was read at the 10th All-Union Conference on Nuclear Spectroscopy, which was held in Moscow from January 19 to January 27, 1960. Report is given in the present paper on the results obtained by experiments, in which the first level of Ne<sup>21</sup> was excited The y emission of graphite, aluminum, and molybdenum targets was studied during their bombardment with Ne<sup>21</sup> ions, whose energy was 24.2 Mev. In all cases a  $\gamma$ -line with 0.35 Mev was found. The authors arrive at the conclusion that these lines, which are known already from previous papers, are not the result of a nuclear reaction but of a Coulomb excitation of  $Ne^{21}$ From publications it is known that the quantum characteristics for the Ne ground state are  $3/2^{+}$ , and for the first excited state  $3/2^{+}$  or  $5/2^{+}$ . Thus, Card 1/2

The Coulomb Excitation of the Ne<sup>21</sup> Rucleus

S/048/60/024/012/007/011

B019/B056

the authors obtained 0.025·10-48e<sup>2</sup>cm<sup>4</sup> or 0.017·10-48e<sup>2</sup>cm<sup>4</sup> for the probability B(E2) of a transition from the ground state to the first excited state. For the partial lifetime of the first excited state one excited state. For the sec or 9.2 10 sec. The authors thank A.B.Girshin thus obtains 6.3·10 sec or 9.2 10 sec. The authors thank A.B.Girshin for the faultless operation of the cyclotron. There are 1 figure and 8 references: 3 Soviet, 4 US. and 1 Danish

ASSOCIATION:

Fiziko-tekhnicheskiy institut Akademii nauk SSSR (Institute of Physics and Technology of the Academy of Sciences USSR) of Physics and Technology of the Academy of Sciences USSR)

s/048/61/025/001/013/031 B029/B060

24.6520 (1138, 1395, 1160)

Andreyev, D. S., Grinberg, A. P., Yerokhina, K. I.,

Lemberg, I. Kh.

Coulomb excitation of the nuclear levels of  $P^{31}$ ,  $S^{33}$ ,  $Mn^{55}$ , TITLE:

and  $Pr^{141}$  by means of  $Ne^{20}$  ions

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, v. 25, PERIODICAL:

no. 1, 1961, 70-76

TEXT: The measurements were made by means of a scintillation spectrometer with a NaI-Tl orystal (40 mm in diameter, 40 mm in height) and by means of a photomultiplier of the type  $\phi 3y - 11$  (FEU-11). The electronics consisted of a pre-amplifier, a cathode follower, an overchargeable amplifier and an AMA-2 (AMA-2) 63-channel pulse height analyzer connected in parallel and an AV 100-1 (AI 100-1) 100-channel pulse height analyzer. Method of measurement, apparatus, and course of the calculation of the reduced transition probability B(E2) have already been

Card 1/7

AUTHORS:

CIA-RDP86-00513R000101510015-7" APPROVED FOR RELEASE: 03/20/2001

Coulomb excitation of the nuclear levels...

S/048/61/025/001/013/031 B029/B060

described in three previous papers (Refs. 1-3). In the case of nitrogen ions the errors are below 15%, but they may attain from 20 to 25% for neon ions. The following was observed when measuring the energy of beam particles by means of deflection in a magnetic field: After deflection, the beam is split into several components corresponding to different charges of the accelerated ions. The change of the ions falling into the beam catcher causes the change of the ratio current strength / number of beam particles in the beam, which means that it influences the accuracy of calculation of the Coulomb excitation cross section. In the experiments concerned, the bombarding particles were quadruple-charged Ne20 ions with energies of 23.2 and 27.8 Mev. The amperage of the ion beam measured on the target was  $\sim 1.10^{-8}a$ . The measurement results are given in the attached Table.  $R_0 = 1.2 \cdot 10^{-13} A^{1/3}$  cm was set. Figs. 1,2, 3,4 show the instrumental y-spectra taken with Ne<sup>20</sup> ions. The following notes are added concerning the individual elements: P31: The Coulomb excitation of the level with  $\Delta E = 1.26$  MeV of P31 was examined with the aid of ions Ne<sup>20</sup> with an energy of 27.8 Mev. The target was pressed from a red phosphorus powder. The spectrum contains a gamma line with

Coulomb excitation of the nuclear levels...

S/048/61/025/001/013/031 B029/B060

Hev in No.20. The value of B(E2)? for the 1.26-Mev level of P31 amounted to 0.011:10-48 e<sup>2</sup> cm<sup>4</sup>. The spins of the two states of P31 are known:

Lo = 1/2 T<sub>f</sub> = 3/2 + S<sup>32</sup>: In the work under consideration, S<sup>33</sup> was excited by quadruple-charged No.20 ions with an energy of 23.2. Mev. The gamma spectrum found contains a line with the energy 0.83±0.01 Mev. Mn.55: The Mn.55 was likewise excited by quadruple-charged No.20 ions with 23.2

Mev. This spectrum contains gamma lines with energies of 0.85 and 0.98

Mev. Pr. 141: Fig. 4 shows the spectrum of gamma rays resulting on the irradiation of prascodymium oxide with quadruple-charged No.20 ions (37.8 Mev). The lifetime of the state with AE = 0.142 Mev amounts to 2:10-9 sec, and its joinal lifetime is 4.5·10-7 sec. The article under consideration is the reproduction of a lecture delivered at the 10th Captuary 19 to 27, 1960. There are 4 figures, 1 table, and 24 references: 2550viet-bloc and 171 non-Soviet-bloc.

Card 3/7

Coulomb excitation of the nuclear levels...

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ASSOCIATION: Fiziko-tekhnicheskiy institut Akademii nauk SSSR (Institute

of Physics and Technology, Academy of Sciences USSR)

Ядро	Δ5, MeV	ΔF°, MeV	<u>п(Е2) †</u> × ×10 <sup>св</sup> , см <sup>4</sup>	7(E2), cen	т, сси	T*, CCH	F
Par Saa	1,26 ± 0,02 0,83 = 0,01	1.264 [5] 0.844±0.006 [6]	0,011 0,0019	4,8·10 <sup>-12</sup> 5,2·10 <sup>-11</sup>	1,0.10-13 <5,2.10-11		9,1 1,0
	$\begin{array}{c} 0.98 \pm 0.01 \\ 0.142 \pm 0.003 \end{array}$	0,839±0,005 [7] 0,983 [8] 0,142 [9]	0,012 0,0036	(5÷13)·10 <sup>-12</sup> 4,3·10 <sup>-7</sup>	_	 2·10 * [10,11]	4,0+5 4,3

Card 4/7

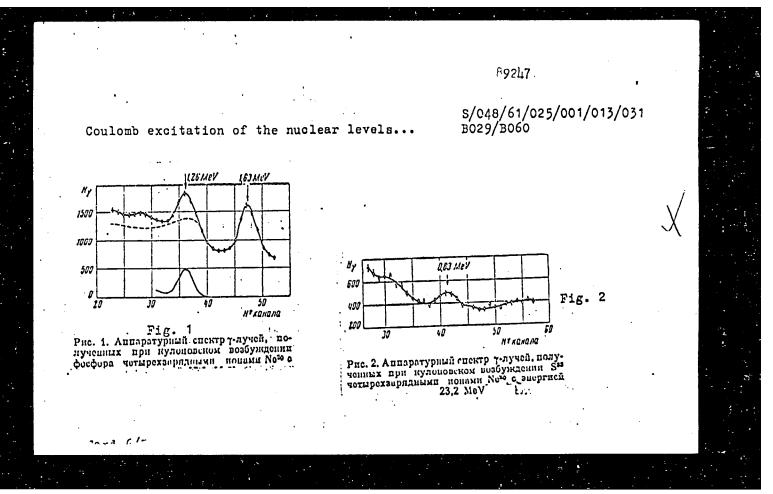
Coulomb excitation of the nuclear levels...

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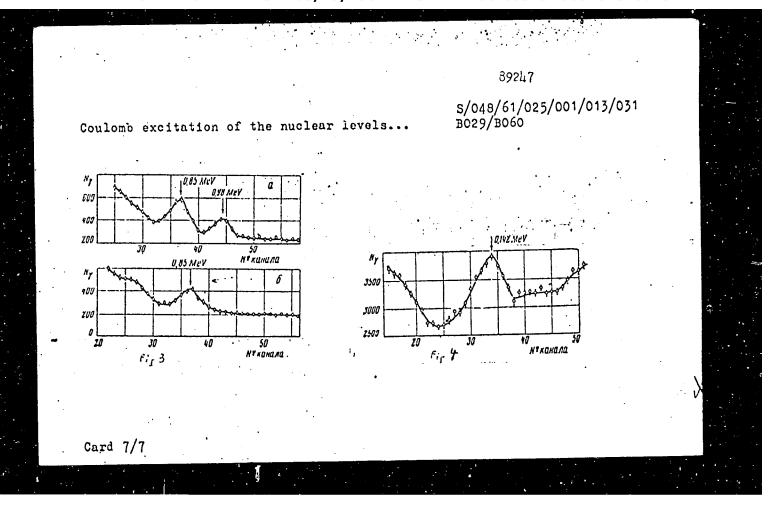
Legend to the Table:  $\Delta E$ , level energy measured in the experiments,  $\Delta E^{*}$ , level energy according to results from other papers,  $B(E2)^{\uparrow}$ , reduced probability of the electric quadrupole excitation of the level,  $\tau(E2)$ , partial lifetime of the level relative to the electric quadrupole transition of the nucleus into the ground state,  $\tau$ , total lifetime of the level,  $\tau^{*}$ , lifetime of the level according to data from other papers, F, ratio between value of  $B(E2)^{\uparrow}$  measured in the experiments and value  $B(E2)_{single\ particle}$  calculated on the basis of the single-particle approximation.

Legend to the Figures: Instrumental  $\gamma$ -spectra, obtained in the case of Coulomb excitation with quadruple-charged Ne<sup>2O</sup> ions of phosphorus at 27.8 Mev (Fig. 1), of S<sup>3O</sup> at 23.2 Mev (Fig. 2); of Mn at 23.2 Mev (Fig. 3)-a without, b with lead filter, 1.25 mm thick; of Pr at 27.8 Mev (Fig. 4).

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24,6300

Card 1/6

Andreyev, D. S., Vasil'yev, V. D., Gusinskiy, G. M., AUTHORS:

Yerokhina, K. I., and Lemberg, I. Kh.

Study of the Coulomb excitation of nuclear levels with the TITLE:

aid of accelerated multiply charged ions

PERIODICAL: Akademiya nauk SSSR. Izvestiya. Seriya fizicheskaya, v. 25,

no. 7, 1961, 832 - 847

TEXT: This paper was read at the XI Annual Conference on Nuclear Spectroscopy in Riga, January 25 - February 2, 1961. The purpose of the studies was to improve the results of earlier work (Ref. 1: Andreyev, D.S. et al., Nucl. Phys., 19, 400 (1960); Ref. 2: Alkhazov, D. G. et al., Zh. eksperim. i teor. fiz., 37, 1530 (1959)) by the method of reference levels (Ref. 1) which consists in measuring the excitation cross section of a reference level before and after measuring the excitation cross section of the level to be investigated. The following nuclear levels were used as reference levels: 0.44~MeV of Na<sup>23</sup> (B(E2)↑ =  $0.11 \cdot 10^{-48}~\text{e}^2\text{cm}^4$ ) for studying Li and B; 1.19 Mev of Ni<sup>62</sup> (B(E2) =  $0.085 \cdot 10^{-48} e^2 cm^4$ ) for

**26439** S/048/61/025/007/001/005 B108/B209

Study of the Coulomb ...

studying Co; 0.76 Mev of Se<sup>76</sup> (B(E2)) = 0.42 · 10<sup>-48</sup> e<sup>2</sup>cm<sup>4</sup>) for studying Mg, Ca, and Se; 1.15 Mev of Sn<sup>122</sup> (B(E2)) = 0.26 · 10<sup>-48</sup> e<sup>2</sup>cm<sup>4</sup>) for studying In, Sb, and Ce; 1.60 Mev of Ce<sup>140</sup> ( $\tau = 1.1 \cdot 10^{-13}$  sec) for studying the even Sn isotopes and Ba<sup>138</sup>; 0.16 Mev of Ti<sup>47</sup> (B(E2)) = 0.040 · 10<sup>-48</sup> e<sup>2</sup>cm<sup>4</sup>) for studying Sn<sup>117</sup>. The excitation probability, B(E2), was determined with an error of 15 - 20%. Tables 1 and 2 contain the results of measurements. In all these studies, the authors made use of the broadening of the energy band of multiply charged ions accelerated in the cyclotron at the FTI (Institute of Physics and Technology). Ne ions having 16 - 18 Mev were used for studying the nuclear levels of light elements such as Li and B, and were also successfully applied to exciting higher levels in light and medium elements (Mg<sup>25</sup>, Mg<sup>26</sup>, Ca<sup>44</sup>, Co<sup>59</sup>, In<sup>115</sup>, and Sb). 52.5-Mev ions of N were able to excite the levels with energies of 1.4 - 1.6 Mev of heavier nuclei (Ba<sup>138</sup> and Ce<sup>140</sup>). The nuclear levels of even-even isotopes were chiefly examined to complete the data on even-even nuclei and to compare results (Ref. 16: Kisslinger, Card 2/6

**26439**, \$/048/61/025/007/001/005 3108/3209

Study of the Coulomb ...

L. S., Sorensen, R. A., Dansk. Mat.-Fys. Medd., 32, No. 9 (1960)) (cf. Table 3). There are 16 figures, 3 tables, and 42 references: 7 Soviet-bloc and 31 non-Soviet-bloc.

Table 1. Coulomb excitation of levels (spin 2<sup>+</sup>) in even-even nuclei.

Legend: (1) Isotope, (2) level energy, Mev, (3) excitation probability, (4) level lifetime,  $10^{-13}$  sec, (5) ratio of B(E2)↑ to the same quantity as estimated for a one-particle model the nuclear radius in the calculations was assumed to be  $R_0 = 1.2 \cdot 10^{-13} \, A^{1/3} \, \text{cm}$ .

Table 2. Coulomb excitation of levels in nuclei with odd A and in odd-odd  ${\tt B}^{10}$  nuclei.

Legend: (1), (2), (3) see Table 1, (6) nuclear spin in ground state, '(7) nuclear spin in excited state, (8) partial lifetime of the level relatively to the electric quadrupole transition, sec.

Legend to Table 3: (1) Nucleus, (2) calculated value of B(E2) as taken from Ref. 16, (3) experimental value of B(E2). Card  $\frac{3}{6}$ 

-ANDREYRY, D.S.; VASIL'YEV, V.D.; GUSINSKIY, G.M.; YEROKHINA, K.I.;

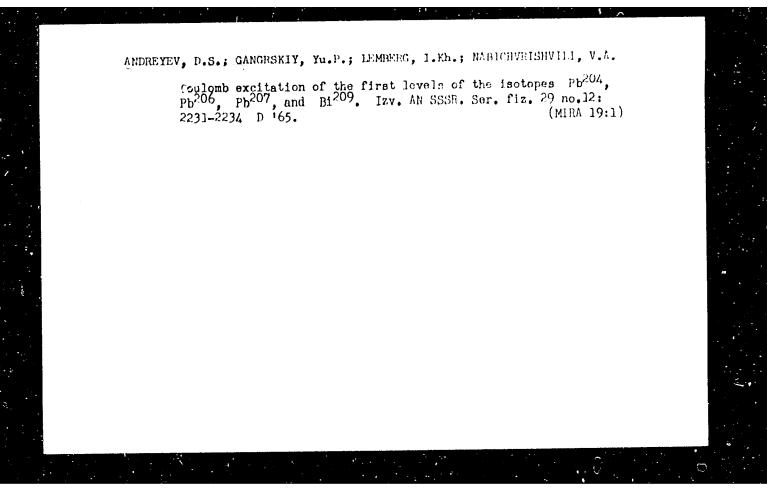
LEMERG, I.Kh.

Investigation of the Coulomb excitation of nuclear levels by the aid of accelerated multiply charged ions. Izv. AN SSSR. Ser. fiz. 25 no.7:832-847 Jl '61. (MIRA 14:7)

(Nuclear reactions) (Ion beams)

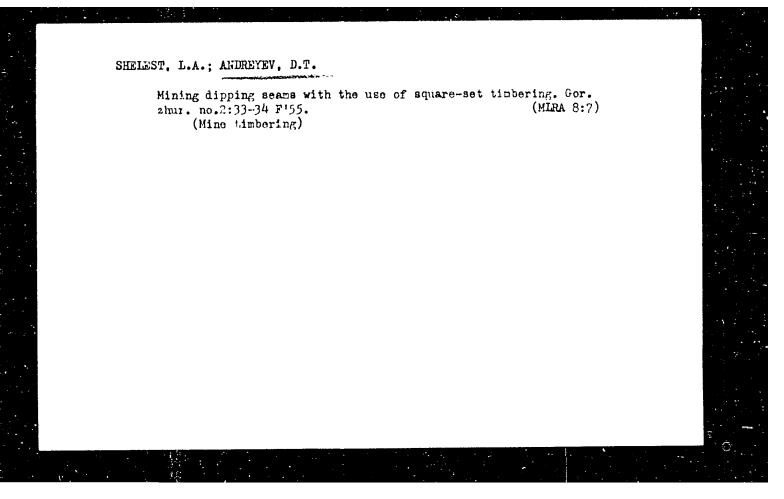
ALKHAZOV, D.G.; ANDREYEV, D.S.; VASIL'YEV, V.D.; GANGRSKIY, Yu.P.; LFMBERG. I.Kh.; VDRALOV, Yu.I.

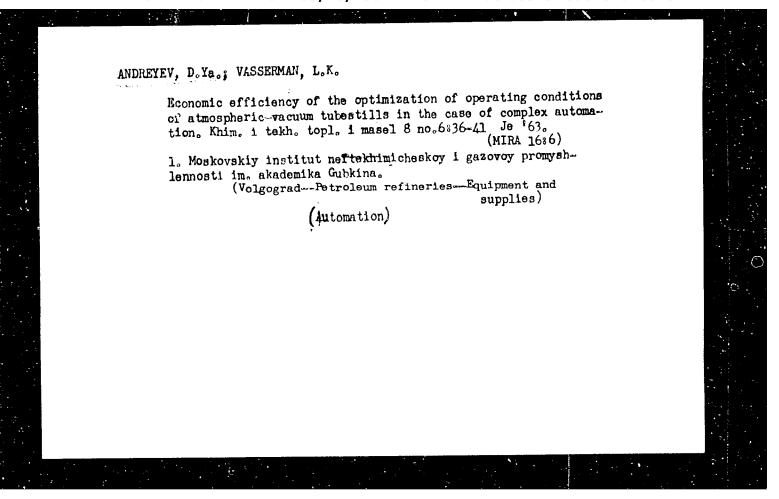
Studying the Coulomb excitation of the first levels of eveneven nuclei by measuring coincidences of gamma quanta and inelastically scattered ions. Izv. AN SSSR. Ser. fiz. 27 no.10:1285-1296 0 '63. (MIRA 16:10)

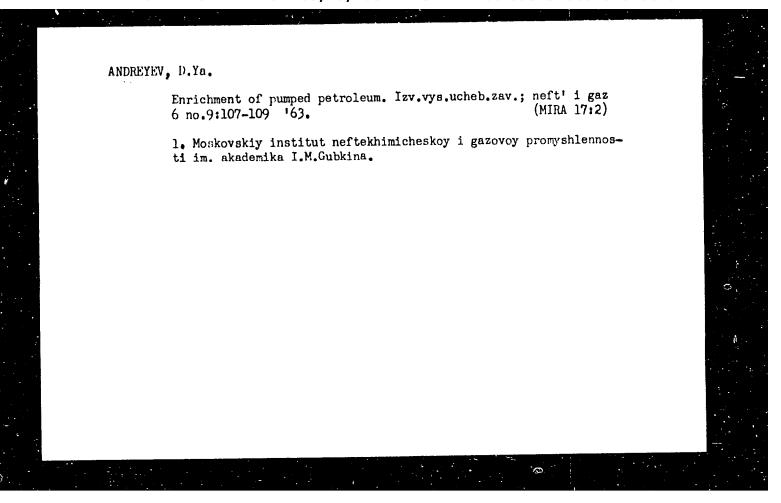


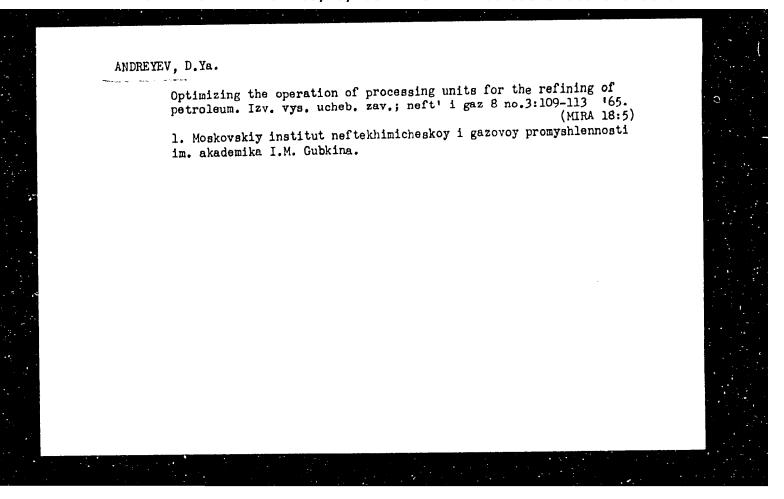
L 26655-66 EWT(m) DIAAP ACC NR: AP6017118 SOURCE CODE: UR/0048/65/029/012/2231/2234 AUTHOR: Andreyev, D. S.; Gangrskiy, Yu. P.; Lemberg, I. Kh.; Nabichvrishvili, V. A. ORG: none TITIE: Coulomb excitations of lower levels in the isotopes Pb sup 204, sup 206, sup 207 and Bi sup 209 This paper was presented at the 15th Annual Conference on Nuclear Spectroscopy and the Structure of the Atomic Nucleus, held in Minsk from 25 January to 2 February 1965/ SOURCE: AN SSSR. Izvestiya. Seriya fizioheskaya, v. 29, no. 12, 1965, 2231-2234 TOPIC TAGS: Coulomb excitation, lead, bismuth, mucleon, nitrogen cyclotron, magnetic field, gamma quantum, even nucleus, neutron proton ABSTRACT: In order to determine the effective nucleon charge it is especially important to know transition probabilities for nuclei having one nucleon (or one hole) above the filled shell. Accurate data on this problem are lacking because the Coulomb output of the excited levels of such nuclei are very small even when bombardment particle energies are very high. Nitrogen ions (N14.5+) were accelerated in the FII cyclotron up to 66.5 Mev by enhancing the magnetic field. Gamma quanta ejected forward from a Bi-enriched lead target were recorded. Nitrogen ions of 66.5 Mev energy were used to study the Coulomb excitation of Pb207 and Bi; and 63 Mev ions, for Pb204 and Pb206. Spectra of yN-coincidence are shown in figures and the results of

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other author	rs. The va	lues of B()	E2) are advers	selv affected	by inadequate			
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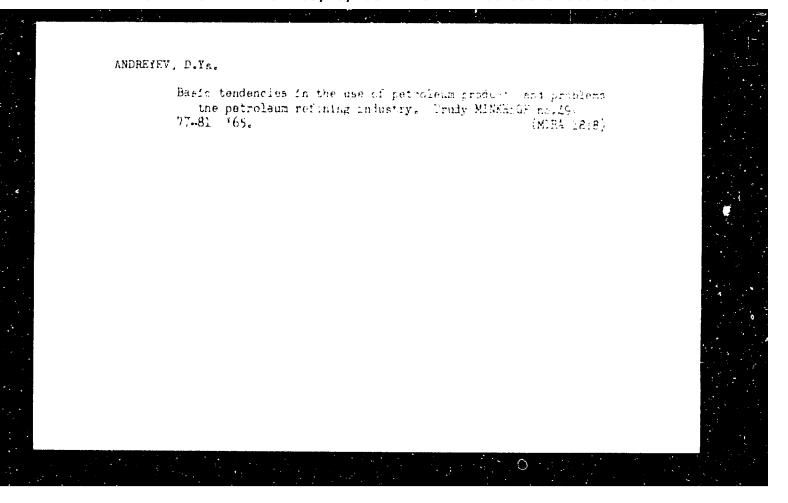




AMDREYEV P.Ya.: Searns, A.D.; Volkeva, I.I., Makindeffi, M.T.

E. m. d. offectiveness of vapital interment in the graduation, gathering, and refinement of potrelear gas. Car. colors. 1, 168-31 (65.)

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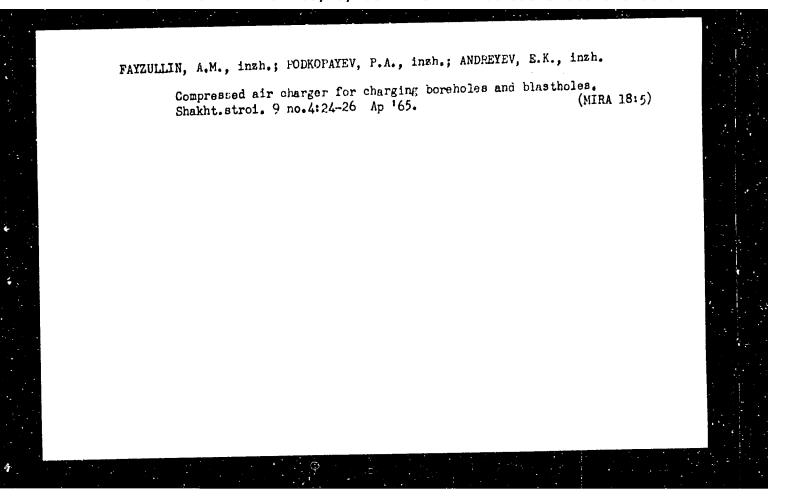


Conducting demonstration experiments in the study of the chemical proporties of oxygen. Khim.v shkole 14 no.3:93
My-Je '59. (MIRA 12:9)

1. Srednyaya shkola pos. Fushkinskiye Gory Pskovskoy obl. (Chemistry -- Study and teaching) (Oxygen)

FAYZULLIN, A.M., ingh.; FODEOFAYEV, P.A., ingh.; ANDERYEV, E.K., ingh.
inequatic ejector charger for charging heles with indunite. Gor.gnur.
no.1:45-47 Ju 165.

(MIRA 18:3)



ACC NR. AR6033921 SOURCE CODE: UR/0196/66/000/007/1003/1003

AUTHOR: Ignatov, V. A.; Andreyev, E. V.

TITLE: Some problems in the designing of end-type induction motors using printed windings

SOURCE: Ref. zh. Elektrotekhnika i energetika, Abs. 7114

REF SOURCE: Tr. Vses. zaochn. energ. in-ta, vyp. 29, 1965, 126-135

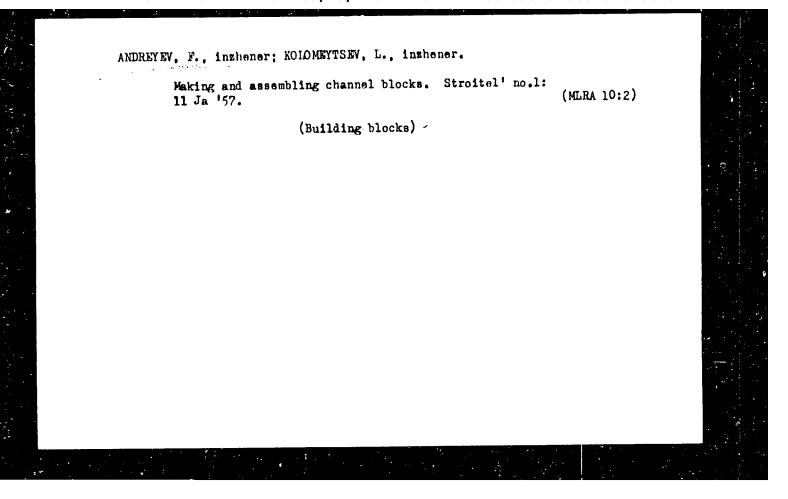
TOPIC TAGS: electric motor, conductor, printed winding

ABSTRACT: A method is given for calculating the dimensions of the printed winding of an end-type electric motor with evolute shaped front sections. The external and internal diameters of the plate, the number of conductors n in the winding, the minimal width of conductor b<sub>min</sub> and the minimal distance between two adjacent conductors b min are given in the calculation. First, the diameter of the base circumference of the front section evolute is determined by the formula

n(b<sub>min</sub> + b min)

Card 1/2

UDC: 621. 3. 045. 049. 75



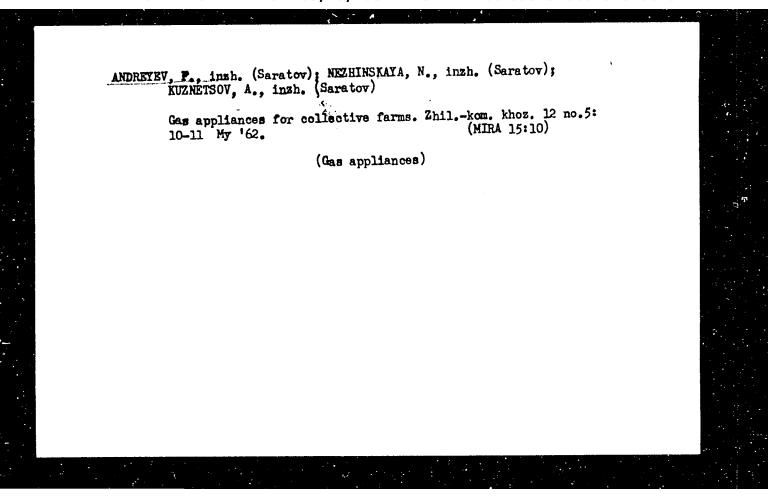
ANDREYEV, F. (g.Saratov); TSENTSIPER, I. (g.Saratov); BOKOVA, Ye. (g.Saratov)

Machine for transporting cylinders of liquid gas. Zhil.-kom. khoz.

11 no.2126-27 F \*61.

(Liquified petroleum gas-Transportation)

(Liquified petroleum gas-Transportation)



ANDROY W, F. A.

Geometry - Study and Teaching

Development of logical thinking in students and solution of proof problems in the 6th and 7th grades. Mat. v shkole No. 1, 1953.

Monthly List of Russian Accessions, Library of Congress, June 1953. Uncl.

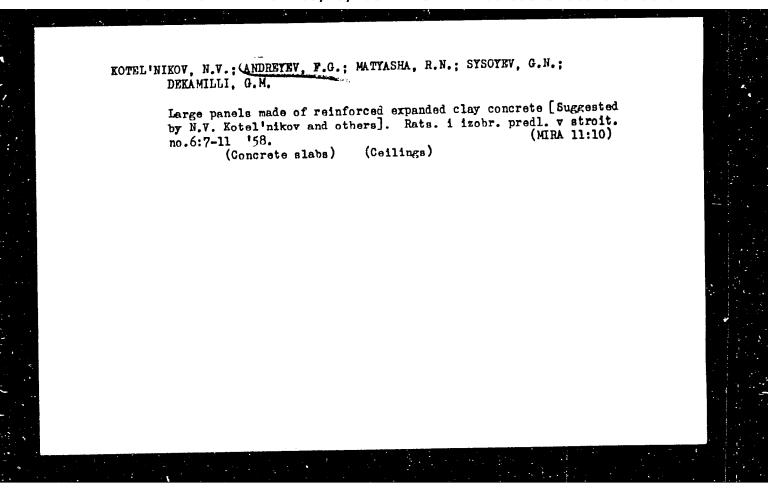
KOTEL'NIKOV, N.V.; MATYASI, R.H.; ANDREYNV, F.G.; SERBYUKOV, N.P.

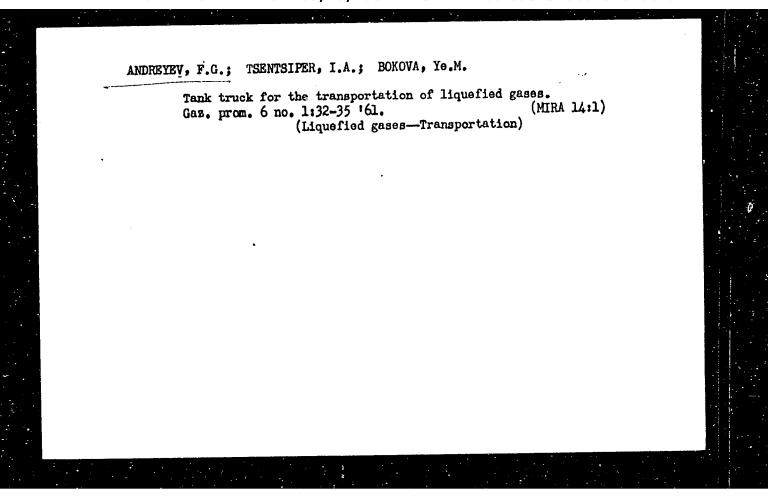
Making concrete wall blocks with flues in construction yards

[Suggested by N.V. Kotel'nikov and others]. Rats. i izobr.

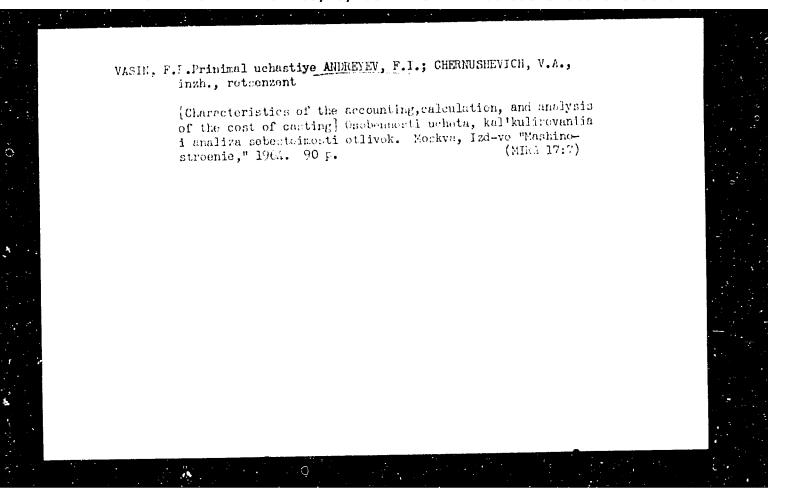
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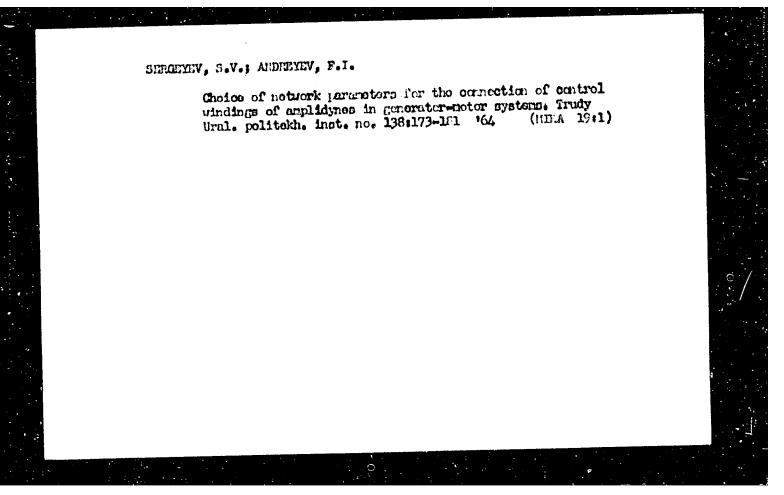
(Concrete blocks)





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TSVETKOV, V.N., kand.tekhn.nauk dots.; ANDREYEV, F.S., inzh.

Construction parameters for fastening porous rubber soles with nylon threads. Izv.vys.ucheb.zav.; tekh.leg.prom. no.5:85-94 (MIRA 13:4)

159.

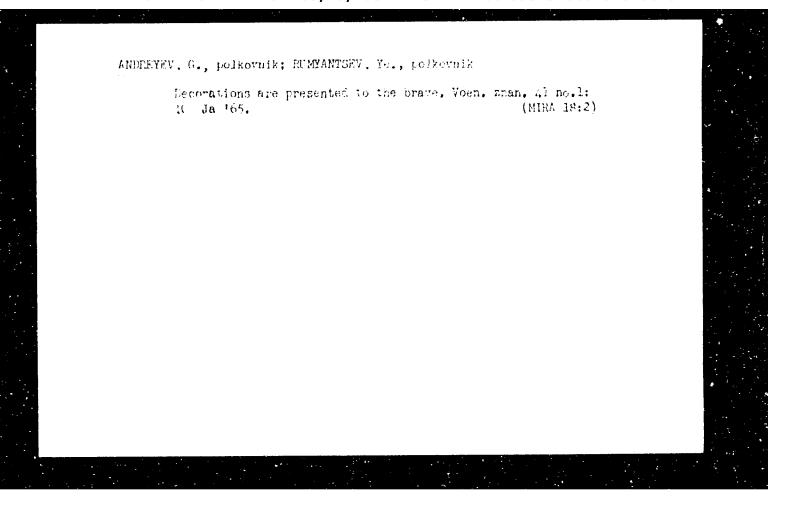
1. Moskovskiy tekhnologicheskiy institut legkoy promyshlennosti.
Rekomendovana kafedroy tekhnologii obuvnogo proizvodstva.

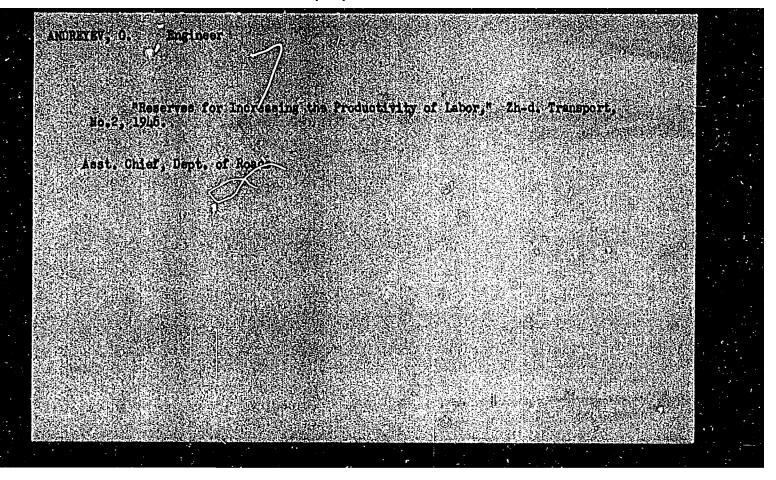
(Boots and shoes, Rubber) (Mylon)

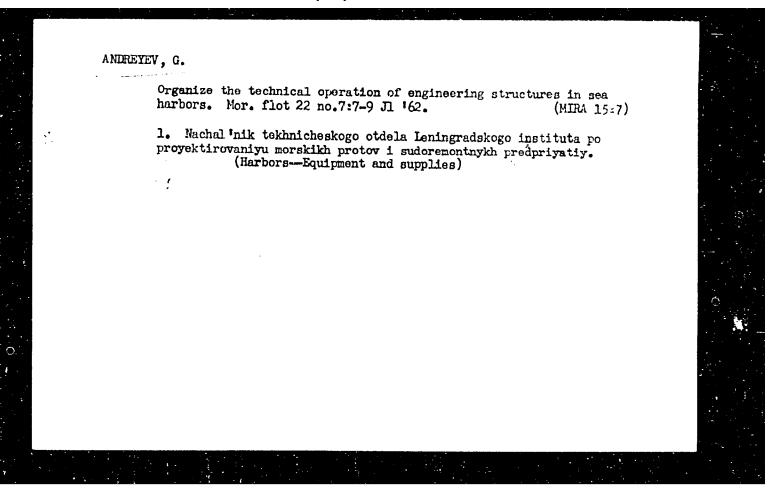
TSVETKOV, V.N., kand.tekhn.nauk, dotsent; ANDREYEV, F.S., inzh.

Efficient parameters for the fastening of porous rubber soles with nylon threads. Report No.2. Izv.vys.ucheb.zav.; tekh.leg.prom. no.1:85-96 160. (MIRA 14:5)

1. Moskovskiy tekhnologicheskiy institut legkoy promyshlennosti. Rekomendovana kafedroy tekhnologii obuvnogo proizvodstva. (Boots and shoes, Rubber)





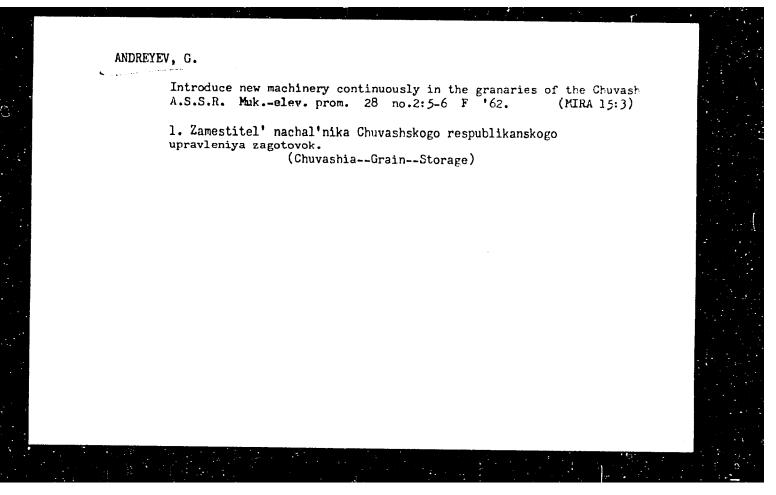


Andreyev, G. - "On the 'terminology and concepts of soil science' by the Academician Prasolov," (In connection with the article of L. I. Prasolov entitled 'Words and Conceptions of Soil Science' published in 'Pochvovedeniye' 1949, No 1), Selektsiya 1 semenovedstvo, 1949, No 6, p. 72-79

SO: U-5240, 17, Dec. 53, (Letopis 'Zhurnal 'nykh Statey, No. 25, 1949).

- 1. ANDREYEV, G.
- 2. USSR (600)
- 4. Cotton Gins and Ginning
- 7. Double-level auger-type gin. Khlopkovodstvo no 8, 1952

9. Monthly List of Russian Accessions, Library of Congress, January 1953, Unclassified.



GAN'ZHIN, V.; FILIPPOVICH, B.; ANDREYEV, G.

Problems in the management and organization of work at grain receiving enterprises. Muk.-elev. prom. 28 no.8:20-22 Ag '62. (MIRA 17:2)

1. Nachal'nik proizvodstvenno-tekhnicheskogo otdela Orenburgskogo upravleniya khleboproduktov (for Gan'zhin). 2. Glavnyy agronom Petropavlovskogo elevatora (for Filippovich). 3. Zamestitel' direktora po kachestvu Tankerisskogo khlebopriyemnogo punkta TSelinogradskoy oblasti (for Andreyev).

ANDREYSV, G., agronom no markshite curtently (Sirgeroderly reyen, Poltavskoy oblasti); SORMAN, D.

More on packaging. Zanbah, rast, et vred. 1 kd. 40 no.30116 '65.

(MRA 18:12)

1. Starshiy agronom pe zantshits rasteviy fractive go rayonnego oblys i nemiya "Sol'khomickingan" (for deflate).